New Platinum Nanocrystal-Based Silicon Nanotubes for Targeting Breast Cancer

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I. Introduction

Cancer is a major cause of death worldwide, and every year millions of people are diagnosed with it. Platinum compounds play an important role as anticancer agents. Their ability to bind to DNA in the nucleus (by a process known as intercalation within DNA base pairs) result in DNA damage and cell death.

Our research group has developed a straightforward method to produce a well-defined nanoscale drug carrier known as silicon nanotubes (SINTs), along with a way to incorporate platinum on their surface using (3-Aminopropyl) triethoxysilane (APTES) as a functional arm. These silicon nanotubes have attracted great attention in applications relevant to diagnosis and therapy, owing in part to its biocompatibility and biodegradability in cells.

II. Methods

A. Synthesis of Silicon Nanotubes (SiNTs)

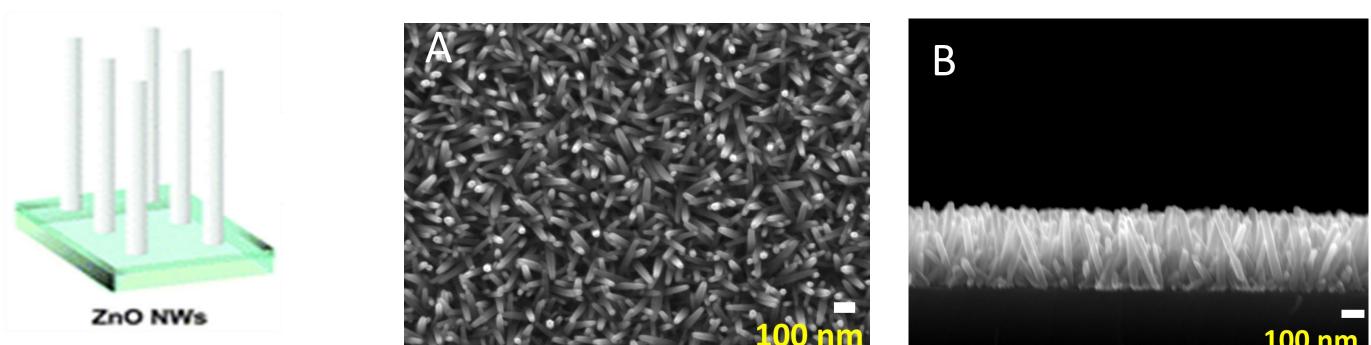


Fig 1. (A) SEM image of ZnO NWs on fluorine-doped tin oxide (FTO) substrate; (B) SEM image cross-section of ZnO NWs on FTO substrate. Growth of ZnO NWs at 95°C.

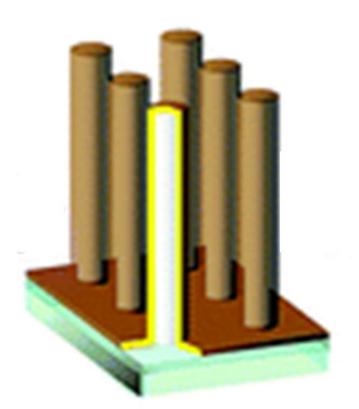
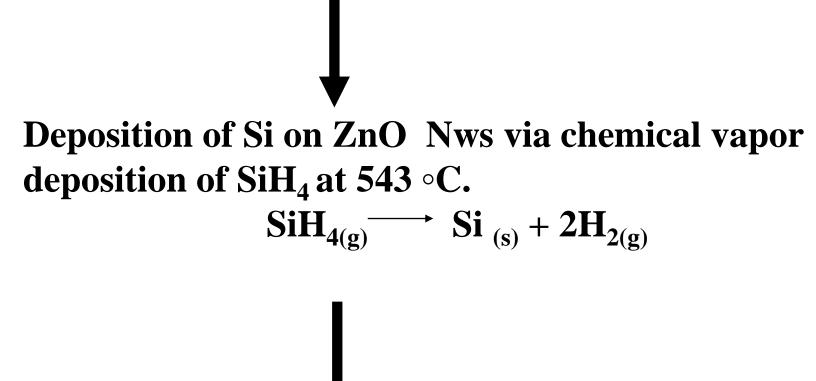


Fig 2. Si/ZnO NWs



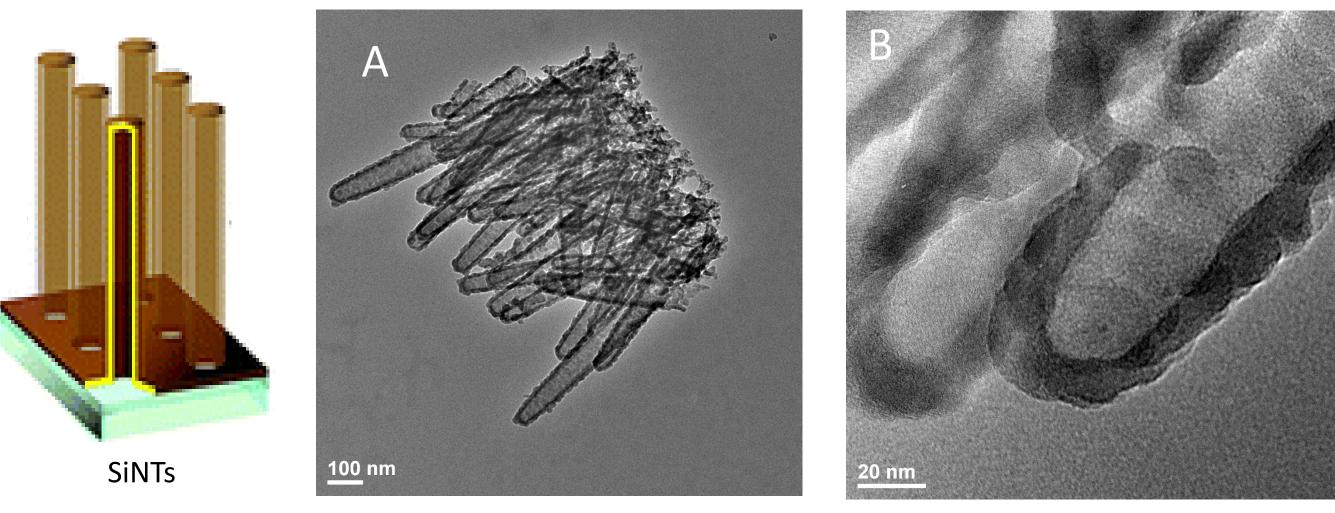


Fig 3 (A) TEM image Silicon Nanotubes (SiNTs) low magnification; (B) Higher magnification

Removal of ZnO NWs to produce hollow SiNTs

 $ZnO_{(s)} + 2HCl_{(g)} \rightarrow ZnCl_{2(s)} + H_2O_{(g)}$ (1) $\operatorname{ZnCl}_{2(s)} + \operatorname{NH}_{3(g)} \rightarrow \operatorname{Zn}(\operatorname{NH}_2)\operatorname{Cl}_{(g)} + \operatorname{HCl}_{(g)}$ (2)



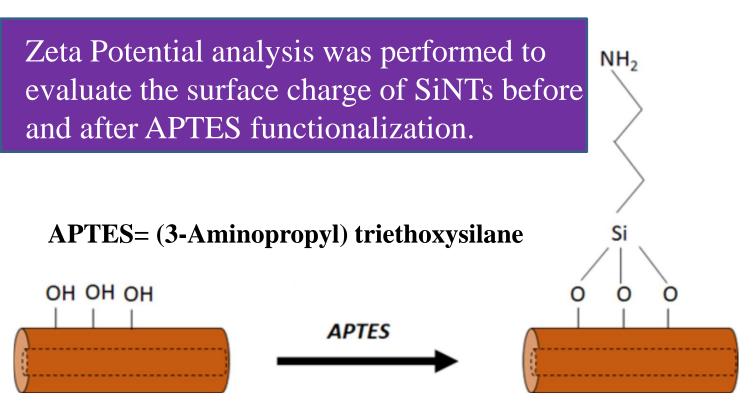
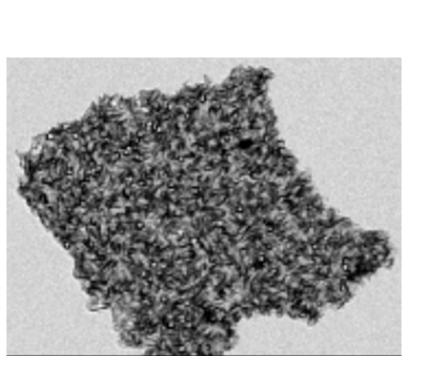


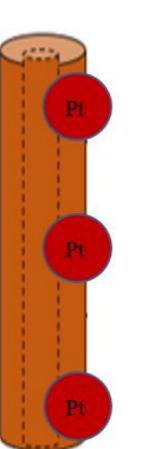
Fig 4. Scheme of functionalization of SiNTs with 2% of APTES in Toluene for 4 h at room temperature

2. Incubation of APTES-SiNTs in K_2PtCl_4 solution at room temperature.

Incubation of APTES-SiNTs was for 24 h 1.5 mM K₂PtCl₄ solution at room Temperature







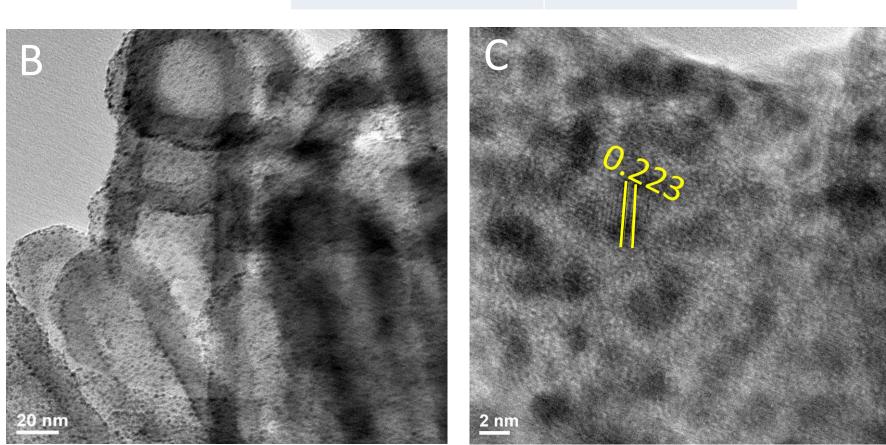


Fig 5. (A) Scheme of platination of APTES-SiNTs with K₂PtCl₄; (B) TEM image of PtNCs-SiNTs low magnification; (C) TEM image of PtNCs-SiNTs highlining the lattice spacing lines.

III. Results

A. Evaluation of Cell Viability of HeLa Cells after Treatment with Pt NCs-pSiNTs (Dr. Nguyen Le.)

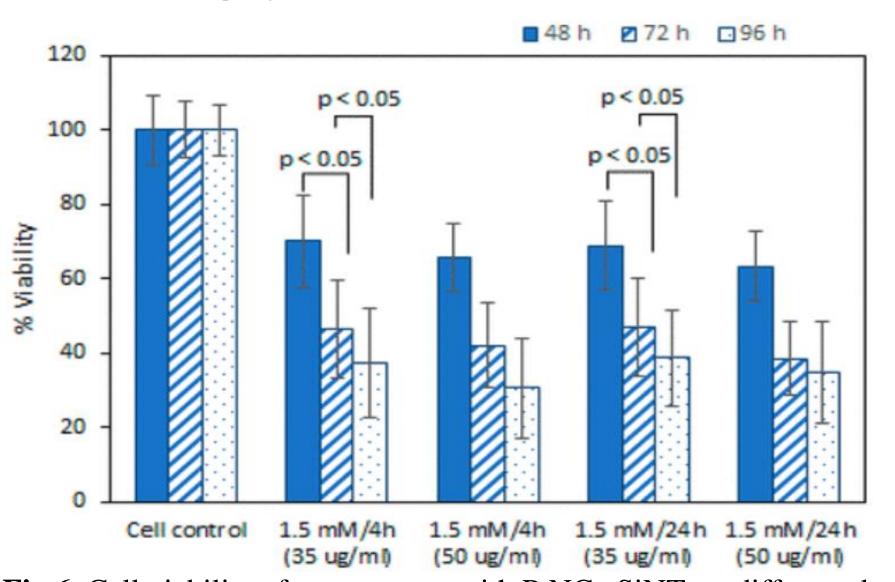


Fig 6. Cell viability after treatment with PtNCs-SiNTs at different doses (2)

 Table 1. Zeta Potential of SiNTs before
 and after functionalization in DI water at pH 6.5

Samples	Zeta Potential (mV)
Unmodified SiNTs	-17.04 ± 0.70
APTES- functionalized SiNTs	+35.87 ± 1.96

 Table 2. Elemental analysis of Bundle
 PtNCs-SiNTs (TEM-EDX)

Element	Wt%
Oxygen (O)	10.64
Silicon (Si)	36.80
Platinum (Pt)	51.64
Chlorine (Cl)	0.93

the molecule-drug

endosome

Endocytic vesicle

C. Conjugation of PtNCs-SiNTs with Folate

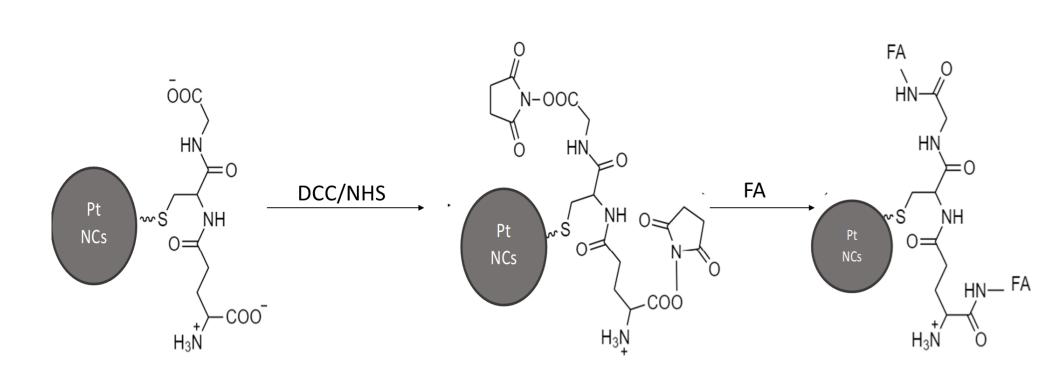


Fig 8. Scheme of Folic Acid attachment with PtNCs
Table 3. Zeta Potential in DI water at pH 6.5

Samples	Zeta Potential (mv)
PtNCs-SiNTs	-28.05 ± 0.67
GSH-PtNCs-SiNTs	-42.38 ± 2.22
Folate-GHS-PtNCs-SiNTs	-18.45 ± 0.93
1 01au-0115-1 UNC5-511115	$-10.4J \pm 0.7J$

IV. Conclusions and Future Work

- attachment with folate

V. References

- *Chem. Commun.*, **2013**,49, 5760-5762

Acknowledgments

- Department of Chemistry and Biochemistry
- Dr. Coffer Group Research



B. Folic Acid as a promising strategy in carcer treatment to direct

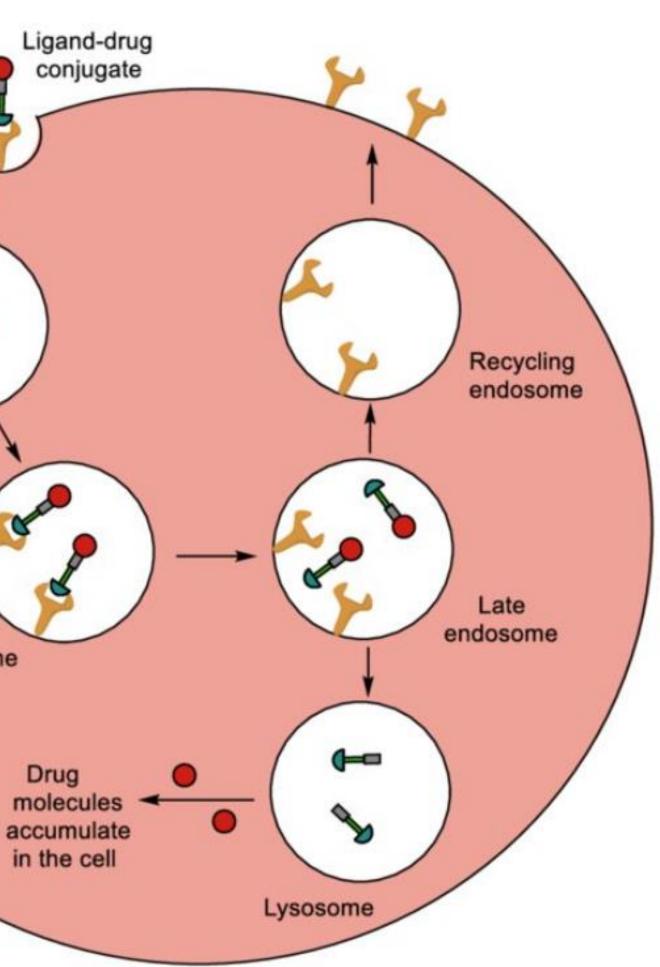


Fig 7. Receptor-mediated endocytosis of a Folate drug-conjugate (3)

• The properties of this material have a high toxicity against HeLa cells inducing apoptosis.

• Results of Zeta potential indicate an attachment of Folate to the PtNCs.

• The next step is performing the X-ray Photoelectron Spectroscopy (XPS) to confirm the

1. Xuezhen Huang, Roberto Gonzalez-Rodriguez, Ryan Rich, Zygmunt Gryczynski and Jeffery L. Coffer.

2. Nguyen T. Le; , Giridhar R. Akkaraju and Jeffery L. Coffer. ACS Appl. Bio Mater. 2020, 3, 1, 208–216 3. Marcos Fernandez; Faiza Javaid and Vijay Chudasama.Chem Sci. 2018 Jan 28; 9(4): 790–810